09994,657

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Search Results - Record(s) 1 through 10 of 10 returned.

[1. 6520950]. 08 May 00; 18 Feb 03. Method of electroporation-enhanced delivery of active agents. Hofmann; Gunter A., et al. 604/503;. A61M025/00.
☐ 2. <u>6399861</u> . 23 May 95; 04 Jun 02. Methods and compositions for the production of stably transformed, fertile monocot plants and cells thereof. Anderson; Paul C., et al. 800/320.1; 800/275 800/288 800/293 800/301 800/302 800/303. A01H005/00 C12N005/04.
3. <u>6350934</u> . 12 Jul 96; 26 Feb 02. Nucleic acid encoding delta-9 desaturase. Zwick; Michael G., et al. 800/281; 435/320.1 435/412 435/419 435/469 435/470 536/23.2 536/23.6 800/278 800/286 800/287 800/292 800/293 800/294 800/300 800/320.1. C12N005/04 C12N015/29 C12N015/82 A01H005/00.
4. 6335161. 25 Feb 98; 01 Jan 02. Release of intracellular material and the production therefrom of single stranded nucleic acid. Martin; Sophie E.V., et al. 435/6; 435/91.2 436/94. C12Q001/68 C12P019/34 G01N033/48.
5. 6329574 . 24 Jul 98; 11 Dec 01. High lysine fertile transgenic corn plants. Lundquist; Ronald C., et al. 800/300.1; 800/278 800/287 800/288 800/293 800/320.1. C12N015/00 A01H001/06 A01H004/00.
6. 6302874. 13 Jul 99; 16 Oct 01. Method and apparatus for electrically assisted topical delivery of agents for cosmetic applications. Zhang; Lei, et al. 604/522; 604/501. A61M031/00.
7. 6103235. 15 Apr 97; 15 Aug 00. Methods of inducing immune tolerance using immunotoxins. Neville; David M., et al. 424/183.1; 424/184.1. A61K039/395 A61K039/00.
8. 6025545. 15 May 95; 15 Feb 00. Methods and compositions for the production of stably transformed, fertile monocot plants and cells thereof. Lundquist; Ronald C., et al. 800/300.1; 536/23.1 536/24.1 800/298 800/300 800/320.1. A01H001/06 A01H004/00 C12M015/00.
9. <u>5990390</u> . 27 Mar 96; 23 Nov 99. Methods and compositions for the production of stably transformed, fertile monocot plants and cells thereof. Lundquist; Ronald C., et al. 800/302; 536/23.71 800/265 800/268 800/320.1. A01H005/00 A01H004/00 A01H001/20 C12H005/04.
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MEDLINE ANSWER 1 OF 4 L2

MEDLINE 90073642 ΑN

DN

A rapid and efficient procedure for transformation of intact Saccharomyces PubMed ID: 2686636 ΤI cerevisiae by electroporation.

ΑU

Department of Biological Chemistry, UCLA School of Medicine 90024. CS

NC

BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, (1989 Nov 15) 164 (3) SO 1157-64.

Journal code: 0372516. ISSN: 0006-291X.

United States CY

Journal; Article; (JOURNAL ARTICLE) DT

English LΑ

Priority Journals FS

198912 EΜ

Entered STN: 19900328 ED Last Updated on STN: 19970203

Entered Medline: 19891228 A rapid and efficient procedure is described for transforming Saccharomyces cerevisiae using electroporation to render intact AΒ cells permeable to DNA. The technique uses relatively low voltages and is particularly sensitive to low concentrations of plasmid DNA. At the highest voltage used (400 volts), the frequency of transformation increased with the amount of plasmid DNA between 25 ng and 100 ng. At higher concentrations of DNA (1-1.5 micrograms) electroporation yielded one-third to one-half the number of transformants obtained with a standard lithium acetate pretreatment. Because this method requires neither pretreatment of cells nor addition of polyethylene glycol (PEG), it has several advantages over currently used transformation procedures.

A rapid and efficient procedure is described for transforming AΒ Saccharomyces cerevisiae using electroporation to render intact cells permeable to DNA. The technique uses relatively low voltages and is particularly sensitive to low concentrations of plasmid DNA. At the highest voltage used (400 volts), the frequency of transformation increased with the amount of plasmid DNA between 25 ng and 100 ng. At higher concentrations.

- ANSWER 2 OF 4 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. L2
- 1990:48268 BIOSIS AN
- BA89:25632 DN
- A RAPID AND EFFICIENT PROCEDURE FOR TRANSFORMATION OF INTACT ΤI SACCHAROMYCES-CEREVISIAE BY ELECTROPORATION.
- SIMON J R; MCENTEE K ΑU
- DEP. BIOLOGICAL CHEM., UCLA SCH. MED., LAB. BIOMED. ENVIRONMENTAL CS SCIENCES, 900 VETERAN AVE., LOS ANGELES, CALIF. 90024.
- BIOCHEM BIOPHYS RES COMMUN, (1989) 164 (3), 1157-1164. SO CODEN: BBRCA9. ISSN: 0006-291X.
- BA; OLD FS
- English LA
- A rapid and efficient procedure is described for transforming AΒ Saccharomyces cerevisiae using electroporation to render intact cells permeable to DNA. The technique uses relatively low voltages and is particularly sensitive to low concentrations of plasmid DNA. At the highest voltage used (400 volts), the frequency of transformation increased with the amount of plasmid DNA between 25 ng and 100 ng. At higher concentrations of DNA (1-1.5 .mu.g) electroporation yielded one-third to one-half the number of transformants obtained with a standard lithium acetate pretreatment. Because this method requires neither pretreatment of cells nor addition of polyethylene glycol (PEG), it has several advantages over currently used transformation procedures.
- A rapid and efficient procedure is described for transforming AB Saccharomyces cerevisiae using electroporation to render intact cells permeable to DNA. The technique uses relatively low voltages and is particularly sensitive to low concentrations of plasmid DNA. At the highest voltage used (400 volts), the frequency of transformation increased with the amount of plasmid DNA between 25 ng and 100 ng. At higher concentrations.
- ANSWER 3 OF 4 CAPLUS COPYRIGHT 2003 ACS L2
- 1976:126439 CAPLUS AN
- 84:126439 DN
- Electrolytic cell for inactivation and destruction of pathogenic material ΤI
- Shaffer, Peter T. B. TN
- Carborundum Co., USA PΑ
- SO U.S., 6 pp.
 - CODEN: USXXAM
- DTPatent
- LA English
- באוז כאוד 1

FAN.	CNT 1 PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	US 3923629 CA 1041038	A A1	19751202 19781024	US 1974-454637 CA 1975-221887	19740325 19750311
PRAI	JP 50133173 US 1974-454637	A2	19751022 19740325	JP 1975-34592	19750324

AB An electrolytic cell for destroying fluid-born

pathogenic materials comprises layers of permeable elec. conductive material sepd. by layers of permeable elec. insulation. conductive layers act as the cell electrodes which are connected to an a.c. source having a current potential of .apprx.0.1-20 volts with frequencies of .apprx.0.1-.apprx.10 Hz. A suitable filter housing surrounds the cell and is arranged so that the pathogen-contg. fluid passes through the permeable electrode layers of the cell. The pathogenic materials are subjected to the elec. potential set up between the layers and are inactivated or destroyed. An electrolytic cell for destroying fluid-born AB pathogenic materials comprises layers of permeable elec. conductive material sepd. by layers of permeable elec. insulation. The conductive layers act as the cell electrodes which are connected to an a.c. source having a current potential of .apprx.0.1-20 volts with frequencies of .apprx.0.1-.apprx.10 Hz. A suitable filter housing surrounds the cell and is arranged so that the pathogen-contg. fluid passes through the permeable electrode layers of the cell. The pathogenic materials are subjected to the elec. potential set up between the layers and are inactivated or destroyed.

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ANSWER 4 OF 4 CAPLUS COPYRIGHT 2003 ACS
L2
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1967:15861 CAPLUS AN

66:15861 DN

Preparation of thoria sols by electrodialysis TI

O'Connor, Thomas L.; Juda, Walter; McNally, Paul H.; Rosenberg, Norman W. IN

APPLICATION NO. DATE

Diamond Alkali Co. PΑ

so U.S., 10 pp. CODEN: USXXAM

DT Patent

LA English

FAN. CNT 1

PATENT NO. KIND DATE US 3280011 19661018 US ______ 19580619 US 3280011 PΙ Fluid, aq. hydrated actinide oxide sols with a controlled and uniform AB particle size, neutral pH, have several advantages over solid reactor fuels. They are prepd. by putting an aq. soln. of a Th metal salt in a 1st chamber of an electrodialysis cell bounded on at least one side by a cation-permeable membrane. In a 2nd chamber on the other side of the membrane H2O is passed. An elec. current is conducted across the membrane through the solns. at room temp. The Th ions in the Th salt soln. pass through the membrane from the 1st to the 2nd chamber forming a Th oxide sol in the 2nd chamber. Thus, a soln. contg. UO2SO4 1, Th(SO4)2 100, H2SO4 5 g., in H2O to make 1 l. in the cathode compartment is subjected to d.c. (100 amp./ft.2 of an electrodialysis cell). The cell consists of 2 chambers sepd. by an anion selective membrane. After 5 hrs. of recirculating electrolysis at 85.degree. and a .apprx.5 volts d.c., the pH increases to 6.2; the actinide sol may then be withdrawn from the compartment. Av. particle size is 55 m.mu. which size is not abrasive to bends and orifices of equipment when pumped at fast rates and not small enough to form gels. Fluid, aq. hydrated actinide oxide sols with a controlled and uniform AB particle size, neutral pH, have several advantages over solid reactor fuels. They are prepd. by putting an aq. soln. of a Th metal salt in a 1st chamber of an electrodialysis cell bounded on at least one side by a cation-permeable membrane. In a 2nd chamber

on the other side of the membrane H2O is passed. An elec. current is conducted across the membrane through the solns. at room temp. The Th ions in the Th salt soln. pass through the membrane from the 1st to the 2nd chamber forming a Th oxide sol in the 2nd chamber. Thus, a soln. contg. U02S04 1, Th(S04)2 100, H2S04 5 g., in H2O to make 1 l. in the

electrodialysis cell). The cell consists of 2 chambers sepd. by an anion

cathode compartment is subjected to d.c. (100 amp./ft.2 of an

selective membrane. After 5 hrs. of recirculating electrolysis at

85.degree. and a .apprx.5 **volts** d.c., the pH increases to 6.2; the actinide sol may then be withdrawn from the compartment. Av. particle size is 55 m.mu. which size is not abrasive to bends and orifices of equipment when pumped at fast rates and not small enough to form gels.

=>



Creation date: 02-03-2004

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